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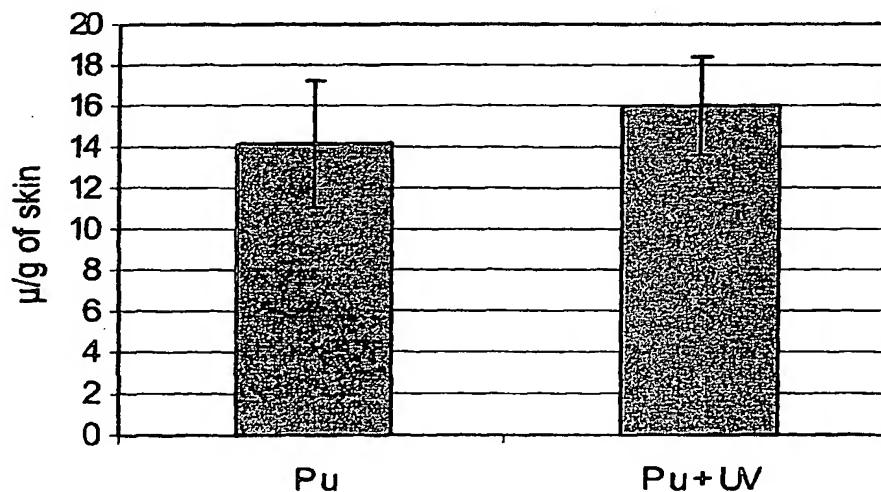
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(54) Title: USE OF POTHOMORPHE UMBELLATA EXTRACT, COMPOSITION ON BASIS OF POTHOMORPHE UMBELLATA EXTRACT AND METHOD OF APPLICATION OF THE POTHOMORPHE UMBELLATA EXTRACT



(57) Abstract: The present invention relates to the use of the Pothomorphe umbellata extract to prepare dermocosmetic and/or pharmaceutical compositions for the treatment and/or prevention of photodamage in the skin, cutaneous aging and/or skin cancer. It also refers to dermocosmetic and/or pharmaceutical compositions for treatment and/or prevention of such photodamage in the skin, cutaneous aging and/or skin cancer prepared on basis of the defined extract. In addition, it provides a method of application of the dermocosmetic and/or pharmaceutical compositions prepared on the basis of the extract of this plant, in way to treat and/or prevent the damage caused to the skin by the excessive exposure to ultraviolet rays of the sun and to artificial tanning lamps.

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USE OF *POTHOMORPHE UMBELLATA* EXTRACT, COMPOSITION ON  
BASIS OF *POTHOMORPHE UMBELLATA* EXTRACT AND METHOD OF  
APPLICATION OF THE *POTHOMORPHE UMBELLATA* EXTRACT

Fields Encompassed by Invention

5       The present invention is applied to Pharmacy and  
Medicine fields and relates to the use of the *Pothomorphe*  
*umbellata* extract in order to prepare dermocosmetic and/or  
pharmaceutical compositions for the treatment and/or  
prevention of photodamage in skin, cutaneous aging and/or  
10 skin cancer. Another form of usage is the application of  
dermocosmetic and/or pharmaceutical compositions on basis  
of *Pothomorphe umbellata* extract, in order to treat and/or  
prevent the damage caused to the skin by the excessive  
exposure to ultraviolet rays of the sun and to artificial  
15 tanning lamps and alterations caused by chronological  
aging.

Background Art

The skin is the human body largest organ and it is  
constantly exposed to the free radicals generating sources,  
20 as ultraviolet radiation, air pollutants and ionizing  
radiation. The multiple exposure to solar radiation,  
without appropriate protection, can produce undesirable  
effects, such as photoaging and photocarcinogenesis  
(TEDESCO, et al., 1997).

25       The aging is followed by a progressive reduction of  
the organic functions and an increase in the vulnerability  
to diseases. Particularly concerning the skin, many  
functional decreases and pathologies connected to peoples  
age are the cumulative result of environmental insults  
30 suffered during intrinsic aging (GILCHREST & YAAR, 1992).  
Intrinsic aging is defined as the changes that occur in all

the individuals throughout time, while photoaging is the overlap of changes attributed to solar exposure throughout the aging process (GILCHREST & YAAR, 1998). Unlike intrinsic aging, not only is the photoaged skin  
5 characterized by an exacerbation of the changes caused by chronological aging, but also for the presence of qualitatively different alterations induced by the exposure to sun (GILCHREST, 1996).

The consequent degenerative alterations of repetitive  
10 exposures to solar radiation occur firstly areas which suffer more exposure, such as face, neck, arms and hands. (DELLA CARBONARE & PATHAK, 1992). Macroscopically, the photoaged skin presents as a dry, nodular surface and with an aspect resembling leather, with deep wrinkles,  
15 accentuated furrows, bags and prominent parts. On the other hand, chronological aging (intrinsic) results in fine wrinkles, tuning and flabbiness of the skin (EMERIT, 1992; RANGARAJAN & ZATZ, 1999). Furthermore, in the photoaged skin, we have the presence of pre-malignant and malignant  
20 neoplasms (TAYLOR et al., 1990).

Those effects seem to be associated with the direct impact of photons in cellular DNA and with the direct impact of free radicals and oxygen reactive species generated from ultraviolet radiation absorption by photo-  
25 sensitive molecules (DELLA CARBONARE & PATHAK, 1992). The consequences of that impact are alteration of the gene expression pattern and the damage to cellular components (SCHARFFETER-KOCHANNEK, 1997). An example is the metaloproteinases synthesis induction by the fibroblasts.  
30 These enzymes are responsible for tissue conjunctive components degradation, such as collagen, elastin,

proteoglycans and fibronectin (SCHARFFETER-KOCHANNEK, 1997; ONISHI et al, 2000).

On the other hand, the skin has a variety of antioxidants, such as enzymatic systems, low molecular weight polar and apolar antioxidants, capable of inhibiting the oxidative damage. The vitamin E is the most important exogenous lipophilic antioxidant found in the tissues. It acts together with antioxidant enzymes, such as glutathione peroxidase, catalase and superoxide dismutase, as well as with smaller molecules as ascorbic acid, glutathione and uric acid. The direct degradation of the  $\alpha$ -tocopherol by UV radiation and the formation of its radicals can influence the other antioxidants of the system, as vitamin C. One of the first events in the cell exposed to ultraviolet radiation is the lipid peroxidation induction. "In vitro" studies, examining low density lipoproteins oxidation, demonstrated that the lipid peroxidation occurs when vitamin E is almost fully degraded. (ESTERBAUER et al., 1993). As aforementioned, the  $\alpha$ -tocopherol indirect degradation occurs by the reaction with peroxy radicals and the formation of oxidized products of  $\alpha$ -tocopherol. Such reactions occur predominantly in the cell lipid core.

There are at least three ways through which the antioxidants concentration can be affected by ultraviolet radiation: (1) direct light absorption, (2) reaction with oxygen reactive species generated by the photosensitization reactions and (3) recovery mechanism in which an antioxidant is saved at the expenses of other (Lopez-Torres et al., 1998).

The chronic oxidative stress, as occurs in modern life due to the excessive exposure to the sun and to the atmospheric pollution increase, may justify a supplementation with antioxidants, in order to delay the cutaneous aging process (EMERIT, 1992).

In this context, the topical application of antioxidants has been considered a promising strategy in the prevention of this oxidative damage to the skin. Considerable pre-clinical (BISSET, 1990; LOPEZ-TORRES ET AL, 1997; FUCHS, 1998) and clinical (MAYER, 1993) data were obtained indicating the potential use of  $\alpha$ -tocopherol in the photodamage prevention to the skin. Besides the  $\alpha$ -tocopherol, other substances of vegetable origin, such as flavonoids and other fenolic compounds, have been proposed for topical application in order to prevent the photodamage to the skin (BONINA et al., 1996; SAIJA et al., 1998), offering advantages, as easy obtaining and smaller production cost.

Other antioxidants have been studied to be used in cosmetics in order to prevent the photodamage. Among them, vitamin C; nitrous acid compounds that control the peroxide formation; zinc and manganese ions that serve as nutrients for native dermis bacteria, accelerating the active substances secretion, such as the enzyme superoxide dismutase and before the  $\alpha$ -tocopherol (FOX, 2000).

Among the employed strategies in the photodamage prevention to the skin, we can point out the use of protective lotions (BISSET et al., 1987), of tretinoin (FISCHER et al., 1998), of alpha-hydroxyacids (GILCHREST, 1996) and antioxidants (FOX, 2000). As aforementioned, the oxygen reactive species are partly responsible for the

damage caused to the skin by the excessive exposure to ultraviolet radiation (FUCHS et al., 1989; SCHARFFETTER-KOSCHANEK, 1997). Therefore, a decrease in the oxygen reactive species load produced by the exposure to radiation  
5 UV represents a very promising strategy of protection against the photodamage.

Nowadays, attention is drawn to the antioxidants of vegetable origin. Antioxidants originating from natural products which offer new treatment possibilities for  
10 diseases mediated by oxidative stress. Some enzymes and high plants secondary compounds were capable of protecting tissues against oxidative stress through the free radicals and oxygen reactive species inhibition or capture. (LARSON, 1988). A "in vitro" study investigated the potential use of  
15 a rosemary extract (*Rosmarinus officinales*) in the photodamage control to the skin (FOX, 2000). The use of flavonoids (quercetin, hesperetin and apigenin) as photoprotecting agents was investigated. Due to the fact that they inhibit the lipid peroxidation of  
20 phosphatidilcholine liposomes exposed to UV and due to the fact that they present a good cutaneous permeation, it is suggested that the topical application of flavonoids might be a good option for the treatment of diseases caused or exacerbated by ultraviolet radiation in the skin (BONINA et  
25 al., 1996).

Brazil has a flora that is extremely rich in medicinal plants with great potential for supplying these antioxidant agents. Among these plants, the family Piperaceae, in particular the "pariparobas" are used thoroughly in popular  
30 medicine. The roots of *Pothomorphe umbellata* (L.) Miq. were

included in Brazilian Pharmacopoeia's first edition (SILVA, 1926).

The *Pothomorphe umbellata* is a plant frequently found in the states of São Paulo, Minas Gerais, Espírito Santo and in the south of Bahia, and popularly known as "pariparoba". This is used in popular medicine for treatment of several illnesses, such as: hepatic insufficiency, indigestion, asthmatic bronchitis and externally in the treatment of burns and common wounds (MORAES, 1983). Brazilian Pharmaceutical Code's first edition depicts the flowing extract of pariparoba, the pariparoba syrup and the pariparoba depurative and ferruginous syrup (MORAES, 1983). The empiric observation of the physiologic action of that plant led to its employment internally and in small doses, as exciting of the stomach and liver functions, by increasing the appetite, activating the digestion such as an aromatic bitterness and promoting the bile drainage, as cholagogue that is. However, such plants are still being used and empirically as antispasmodic, which still has not been confirmed by clinical assays. Internally, these plants effects are recognized in the country, evidently, and have a great merit and acceptance, that comes from the aborigines (FREITAS, 1999; PECKOLT, 1941).

In relation to the antioxidant activity of the lyophilized extract of the *Pothomorphe umbellata* root, in vitro assays have evaluated it, using as model the self-oxidation of mouse brains, and such activity was partly attributed to the presence of 4-nerolidylcatechol (BARROS et al., 1996), a fenolic compound extracted from the vegetable roots (KIJJOA, 1980). In another in vitro assay,



the *Pothomorphe umbellata* roots extract showed an antioxidant potential significantly larger than the one of this isolated compound, suggesting the presence of additional compounds with antioxidant activity (DESMARCHELIER et al., 1997). Based on these data, a study was accomplished seeking to evidence and evaluate the antioxidant activity of the *Pothomorphe umbellata* extract in the skin. The topical application of an lyophilized extract of the *Pothomorphe umbellata* roots on the skin of Hairless mice caused a reduction of 97% in the lipoperoxidation indicators as production of reactive substances to the acid tiobarbituric and chemiluminescence. This antioxidant activity was 2,5 times larger than the one of the  $\alpha$ -tocopherol, a known antioxidant, applied to the same conditions (RÖPKE, 1999).

Although the antioxidant activity of *Pothomorphe umbellata* is known, there are no available information regarding the absorption of the active principle by the skin and about the most appropriate formulation to serve as vehicle for this drug. Furthermore, there are not specific studies about the performance of the active principle of this plant in the oxidative stress caused by the ultraviolet radiation.

#### Objects, Advantages and Solutions

The present invention intends to demonstrate the antioxidant and inhibitory action of the lipoperoxidation in the skin presented by the pariparoba extract (*Pothomorphe umbellata*), the preparation of dermocosmetic and/or pharmaceutical compositions capable of treating and/or preventing the photodamage in the skin caused by the excessive exposure to the sun and to help in the treatment

of photoaging and/or skin cancer, as an alternative to the substances of vegetable origin presented in the state of the art. Also, it proposes an application method of the dermocosmetic and/or pharmaceutical compositions on basis  
5 of the extract of this plant.

The employment of *P. umbellata* extract in the preparation of a medicament is quite interesting for its economic feasibility, due to the easy access to the raw material, as it is very common Brazilian plant with low  
10 production cost, and due the fact that the extract can be obtained in a simple way, not involving complex and costly techniques.

#### Drawings

In the following, the invention will be described and  
15 for a better understanding, figures will be presented:

**Figure 1:** Acid ascorbic concentration ( $\mu\text{g/g}$  of skin) in the skin of mice. The values are represented as percentages of the average acid ascorbic concentrations in the skin of control mice ( $174.61 \pm 7.76 \mu\text{g/g}$  of skin);  
20 percentage of the irradiated control group (C + UV),  $80.063 \pm 14.57$ ; of the gel irradiated control group (G + UV)  $78.59 \pm 19.04$  and of the *P. umbellata* gel treated group (Pu + UV),  $103.74 \pm 22.04$ . The difference among the groups was not considered significant. The values were obtained from  
25 the average of the results of seven animals.

**Figure 2:**  $\alpha$ -tocopherol concentration ( $\mu\text{g/g}$  of skin) in the skin of the mice. The values are represented as percentages of the average  $\alpha$ -tocopherol concentrations in the skin of mice of control group ( $2.70 \pm 0,69 \mu\text{g/g}$  of  
30 skin). Percentage of the irradiated control group (C + UV),

35.05  $\pm$  11.26; of the gel irradiated control group (G + UV) 42.90  $\pm$  20.28 and of the *P. umbellata* gel treated group (Pu + UV), 106.77  $\pm$  28.

\*p <0,001, considered very significant when compared to the control group. The values were obtained from the average of the results of seven animals.

**Figure 3:** 4-nerolidylcatechol concentration in the skin of irradiated and not irradiated mice, treated with *P. umbellata* gel (Pu, 14.14  $\pm$  3.09  $\mu$ g/g of skin) and with radiation UVB (Pu + UV, 16.00  $\pm$  2.38 $\mu$ g/g of skin). The data represent the average and the standard deviation of 6 animals.

**Figure 4:** Hairless mouse skin exposed to ultraviolet radiation (degree 4).

**Figure 5:** Hairless mouse skin exposed to ultraviolet radiation and treated with the gel containing *Pothomorphe umbellata* extract (degree 1).

#### Description of the Invention

In the present invention, the pariparoba (*Pothomorphe umbellata*) extract is extracted from the root of this plant (commonly found in the south-eastern area of Brazil). It presents, among others, antioxidant activity, preventing the photodamage caused by the excessive exposure to the sun and to artificial tanning lamps. It also presents inhibitory action of the lipid peroxidation, which is one of the first events to occur in the cell exposed to ultraviolet radiation.

The skin is equipped with a variety of antioxidants as enzymatic systems, low molecular weight polar and apolar antioxidants, capable of inhibiting the oxidative damage. The  $\alpha$ -tocopherol indirect degradation, that is an

antioxidant, it occurs through the reaction with peroxy radicals and formation of  $\alpha$ -tocopherol oxidated products. These reactions occur predominantly in the cell lipidic medium. One of the performing forms of the *P. umbellata* extract is over those antioxidants in order to prevent the effects caused by the skin oxidative stress, that is, the 4-nerolidylcatechol found in the pariparoba extract acts on the  $\alpha$ -tocopherol in order to avoid its degradation and consequently the one of other antioxidants of the enzymatic system.

The pariparoba (*P. umbellata*) antioxidant action, as mentioned above, is caused partly by the presence of the 4-nerolidylcatechol in the extract of that plant. However, as previously seen, studies (BARROS et al., 1996) showed that the extract, when being used, presented higher activity than the 4-nerolidylcatechol in an isolated way. Therefore, the proposed invention uses the extract obtained from the root of that plant, for the obtaining of dermocosmetic and/or pharmaceutical compositions.

#### 20 Pharmaceutical Composition

Another aspect of the present invention refers to the dermocosmetic and/or pharmaceutical composition, which comprises as an active ingredient, the *P. umbellata* root extract. Composition shall be considered as the group formed by the active principle and the other ingredients (pharmaceutically acceptable excipient) that form the carrier, as well as any product that results, direct or indirectly, from the dissociation of one or more of the ingredients, or of other types of reactions or interactions of one or more of the ingredients.

The composition, exemplified in the form of a dermocosmetic, according to the proposed invention, can be prepared in accordance with prior art methods, for topical use. This example illustrates the chosen formulation, but  
5 it does not intend to limit the invention in any way. A proposed composition is comprised of:

- a) carboxymethylcellulose 0.01 - 10.0%
- b) propyleneglycol 0.001 - 50.0%
- c) methylparaben 0.001 - 3.0%
- 10 d) *Pothomorphe umbellata* standardized extract, so that the formulation contains 0.005 to 20.0% of 4-nerolidylcatechol
- e) distilled water q.s.p. 100.0%

For the preparation of an effective composition, it is  
15 necessary that the extract standardization in regard to the amount of 4-nerolidylcatechol present therein. This is made in a high efficiency chromatography device coupled to an electrochemistry detector or UV detector.

The preparing of the composition of this invention can  
20 be made through any known methods in the pharmacy art and in combination with an pharmaceutical carrier, in accordance with the conventional techniques of pharmaceutical composition. The proposed composition is presented in the form of gel. But the presentation can be  
25 made in several forms, depending on the preparing possibility for the desired topical use, for instance, for the lips, labial protective, for the body, protective lotions for the body, lotion, moisturizers, among others. After standardizing the extract in regard to its  
30 concentration of 4-nerolidylcatechol, then, it is incorporated in the cosmetic and/or pharmaceutical bases,

so that to reach the desired active principle concentration.

In all possibilities, they must be sterile and stables in the manipulation and storage conditions and preserved  
5 against polluting action of microorganisms, as bacteria and fungi.

The carrier, in addition to the mentioned ones in the proposed composition, can be a solvent or a dispersion medium containing, for example: water, ethanol, polyol (for  
10 example, glycerol, propylene glycol and liquid polyethylene glycol), appropriate mixtures and vegetable oils. In addition to aforementioned the ingredients, the described pharmaceutical formulations can include, in an appropriate way, one or more carrier ingredients, such as diluents,  
15 buffers, ligands, surface active agents, thickeners, preservatives (including anti-oxidizers) and similar and inclusion of other substances.

#### Doses

The effective dosage of the used ingredient can vary  
20 depending on the way of presentation of the dermocosmetic and/or pharmaceutical composition, the condition to be treated and the severity of the condition to be treated, so that the amount of the *P. umbellata* extract topically administered is effective. In a general way, it can be said  
25 that the formulation will be effective with the presence of at least 0.1% of 4-nerolidylcatechol.

The capacity of the pariparoba extract and accordingly of the 4-nerolidylcatechol of acting in the treatment and/or prevention to the photodamage to the skin, cutaneous  
30 aging and skin cancer can be illustrated together with the following non-limiting examples and figures.

Example 1 - Effect of the topical application of *Pothomorphe umbellata* root extract on the low molecular weight antioxidants, in the oxidative stress of the skin.

Hairless male mice were used (HRS/J), with approximately 10 weeks of age. They were divided into four groups, each one having 7 animals. The first group received topical treatment with gel base without the presence of *P. umbellata* and the second group with *P. umbellata* gel with concentration of 0.1% of 4-nerolidylcatechol. The third group did not receive any treatment. After two hours, the animals were irradiated by a Philips lamp for 30 min with only one dose of UVB radiation (290-320 nm), 26 cm distant. This irradiation corresponds to  $22.95 \times 10^{-2} \text{ J/cm}^2$ , twice the minimum dose to cause erythema in these animals. The radiation was measured with a UVB sensor. Still, a fourth group of mice did not receive treatment, nor it was irradiated.

After the irradiation, the animals were sacrificed by cervical displacement. The 4-nerolidylcatechol absorbed by the skin and the  $\alpha$ -tocoferol of the sacrificed mice tissues were extracted based on the method proposed by Burton (BURTON AND INGOLD, 1985) and, then, it was made the quantitative analysis of these components through the chromatographic method. Was also quantitatively analyzed the ascorbic acid, through the methodology proposed by Wayner & Burton (WAYNER & BURTON, 1989).

The obtained results are shown in the figures 1 to 3. As shown in the figure 1, no significant alteration was noticed in the levels of ascorbic acid in the skin of irradiated mice, possibly due to the used dose and to the UV  $\lambda$  used. In the figure 2, it is possible to observe that

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topical treatment with *P. umbellata* preserved the levels of  $\alpha$ -tocopherol in the skin of irradiated mice, protecting itself against degradation for the UV radiation. After the irradiation, there was a significant decrease ( $p < 0.01$ ) in the levels of  $\alpha$ -tocopherol in the skin of the mice from the control group irradiated, while in the treated group with *P. umbellata* gel, the  $\alpha$ -tocopherol was preserved ( $p > 0.05$ ).

No Difference was observed among the 4-nerolidylcatechol concentration in the skin of irradiated and not irradiated mice, as shown in figure 3. The similar chemical nature of the  $\alpha$ -tocopherol and of the 4-nerolidylcatechol can increase its interaction, explains the preservation of the levels of  $\alpha$ -tocopherol in the mice treated with the *P. umbellata* extract. In the skin of the animals irradiated with UVB ( $0.3 \text{ J/cm}^2$ ), there was a reduction of approximately 60% in the  $\alpha$ -tocopherol concentration.

**Example 2** - Effect of the topical application of formulation on the basis of the *Porhomorphe umbellata* root extract in the prevention of photoaging caused by the chronic exposure to ultraviolet radiation.

It was used in this example a gel formulation containing the *Porhomorphe umbellata* extract which obtained in the previous tests the best liberation of active principle in the skin. I. UVB Philips 15 W 12 rs 40W was used for irradiation of Hairless mice. The animals were irradiated four times a week, for 10 min, during a period of 22 weeks. The used dose was of  $76.5 \text{ mJ/cm}^2$  (approximately 0.7 times the necessary minimum dose to cause erythema in the animals). The dose was measured in a



radiometer, with a UVB SED 240 sensor. Two hours before the irradiation, the mice were treated with gel containing the *Pothomorphe umbellata* extract, in the concentration of 0.1% of 4-nerolidylcatechol. At the same time, mice without treatment and treated with gel without active principle as control groups were irradiated. A fourth control group was not treated nor irradiated.

The formation of wrinkles was classified according to the proposed score in the experimental model of photoaging presented by Bisset (BISSET et al., 1987).

**Table 1:** Score for evaluation of the wrinkles formation degree in the skin of Hairless mice.

DEGREE	Skin Description
0	Numerous fine grooves covering the back and the sides of the body, that appear and disappear with the movement.
1	All the fine grooves in the back and along the spine disappear. Few superficial and rough wrinkles perpendicular to the head-tail direction, that appear and disappear with the movement.
2	Rough and permanent wrinkles along the body.
3	Deep and permanent wrinkles. Leathery skin with no elasticity.

4	Skin with coriaceous aspect having lesions of dark coloration in the back.
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The mice treated with *P. umbellata* gel (figure 5), visibly did not suffer the consequences of the chronic exposure to the ultraviolet radiation, as much as those which were irradiated without being treated with the *P. umbellata* gel (figure 4). This study proved the effectiveness of the formulation in the prevention of photoaging caused by the chronic exposure to ultraviolet radiation. With regard to the animals of other groups, alterations caused by photoaging were noticed, as such deep and permanent wrinkles, leatheriness and even lesions were visible.

It is being demonstrated through to present invention an easy form to obtain *Pothomorphe umbellata* extract and its effectiveness as an antioxidant and inhibitor agent of the lipid peroxidation in the skin, it makes of formulations containing this extract an important alternative for treatment and/or prevention to the photodamage in the skin, cutaneous photoaging and skin cancer, which are one of the health agents' most serious concerns, due to the increase of the exposure to ultraviolet radiation caused by the decrease of the ozone layer of Earth.

CLAIMS

1. **USE OF *POTHOMORPHE UMBELLATA* EXTRACT** comprising the preparation of dermocosmetic and/or pharmaceutical compositions for treatment and/or prevention of photodamage to skin, cutaneous aging and skin cancer.

2. **USE OF *POTHOMORPHE UMBELLATA* EXTRACT** according to claim 1 in which the obtained extract comprises antioxidant activity.

3. **USE OF *POTHOMORPHE UMBELLATA* EXTRACT** according to any preceding claim in which the obtained extract comprises inhibitory activity of the lipid peroxidation.

4. **COMPOSITION ON BASIS OF *POTHOMORPHE UMBELLATA* EXTRACT FOR TREATMENT AND/OR PREVENTION OF PHOTODAMAGE TO , CUTANEOUS AGING AND/OR SKIN CANCER** according to any preceding claim comprising that the *Pothomorphe umbellata* plant extract to be active component.

5. **COMPOSITION** according to claim 4 comprising a formulation which contains a range from 0.005 to 20.0% of 4-nerolidylcatechol in the *Pothomorphe umbellata*.

6. **COMPOSITION** according to claim 4 or 5 comprising acceptable carriers.

7. **COMPOSITION** according to claim 4, 5 or 6 comprising a composition which is presented in all the forms for topical use.

8. **COMPOSITION** according to claim 7 comprising a composition which is presented in gel form.

9. **COMPOSITION** according to claim 8 comprising:

a) carboxymethylcellulose 0.01 - 10.0%

b) propyleneglycol 0.001 - 50.0%

c) methylparaben 0.001 - 3.0%

d) *Pothomorphe umbellata* standardized extract, so that the formulation comprises from 4-nerolidylcatechol 0.005 to 20.0%

e) distilled water q.s.p. 100.0%

5        10. COMPOSITION according to claim 8 comprising a composition which may be presented in the moisturizer form, lotion form, sun protective, labial protective and cream.

10       11. METHOD OF APPLICATION OF THE DERMOCOSMETIC AND/OR PHARMACEUTICAL COMPOSITION ON BASIS OF *POTHOMORPHE UMBELLATA* EXTRACT FOR TREATMENT AND/OR PREVENTION OF PHOTODAMAGE TO SKIN, CUTANEOUS AGING AND/OR SKIN CANCER comprising a dermocosmetic and pharmaceutical compositions to be topically administered in way to allow a satisfactory therapeutic response.

15       12. METHOD according to claim 11 comprising an antioxidant activity.

13. METHOD according to claim 11 comprising an inhibitory activity of the lipid peroxidation.

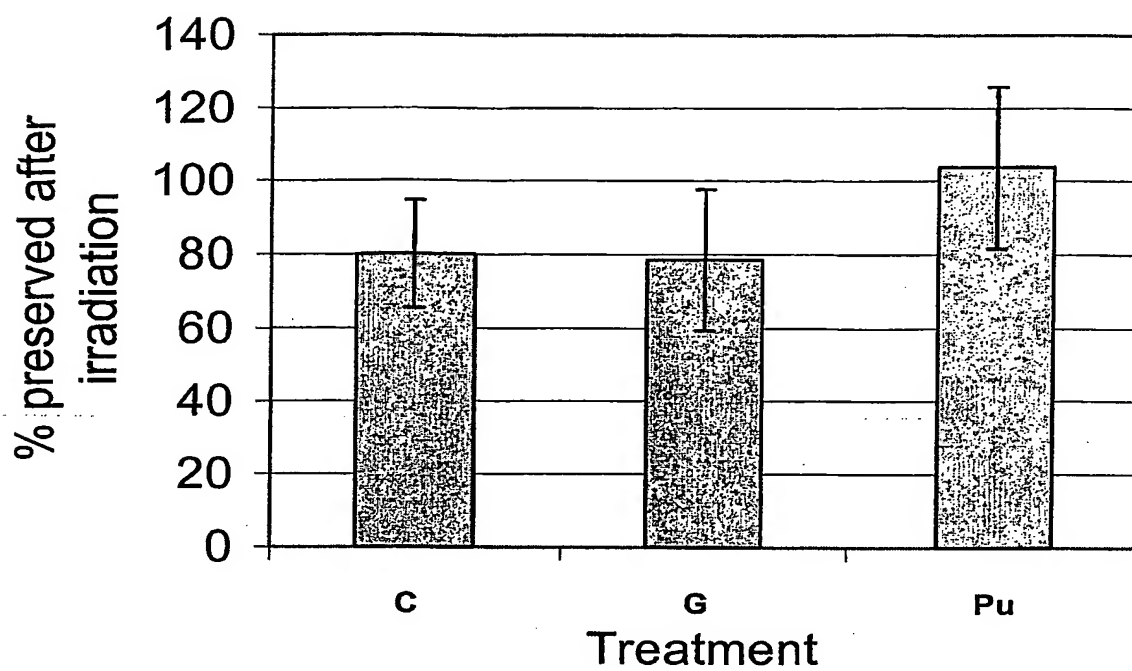


FIGURE 1

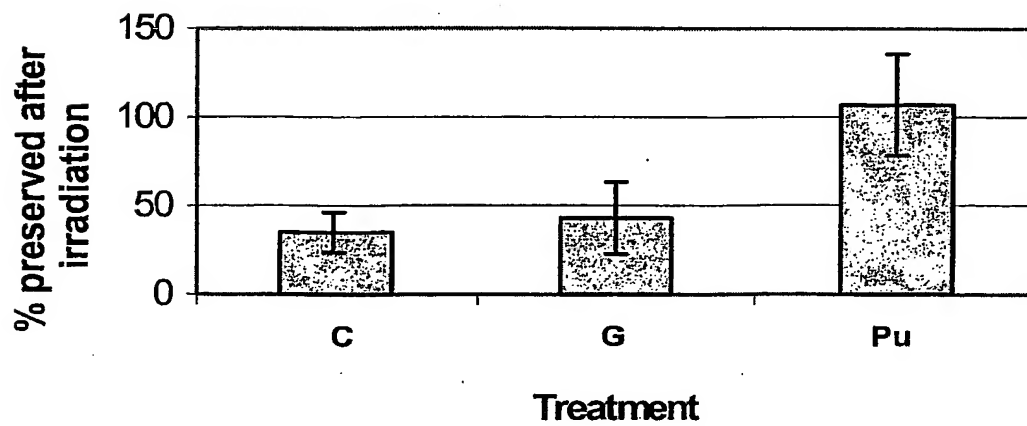


FIGURE 2

2/3

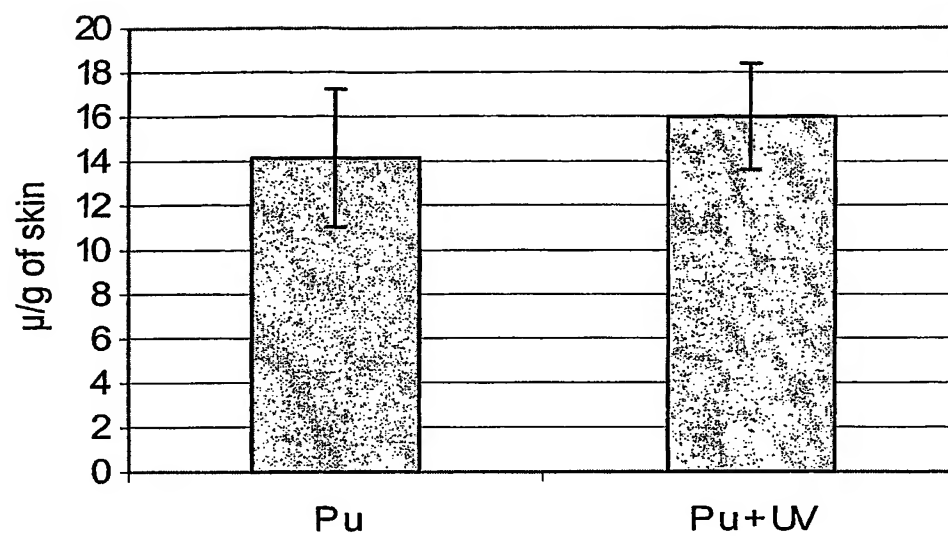
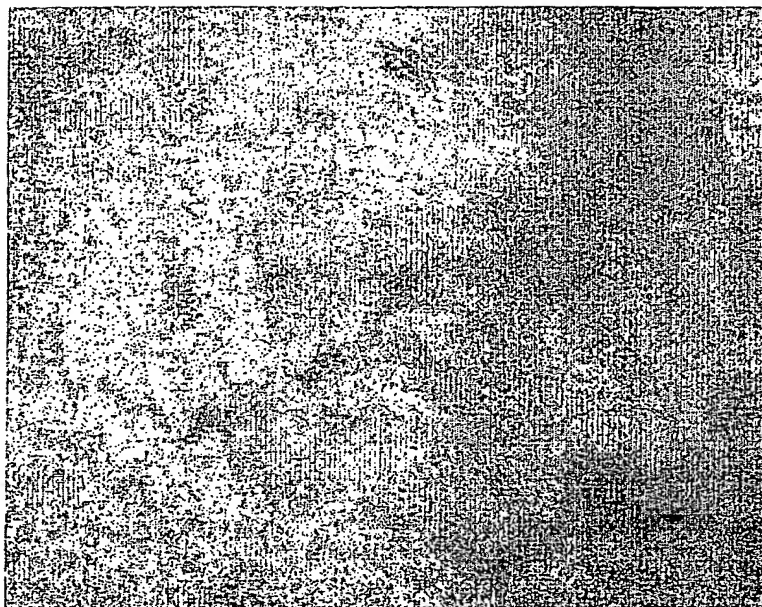
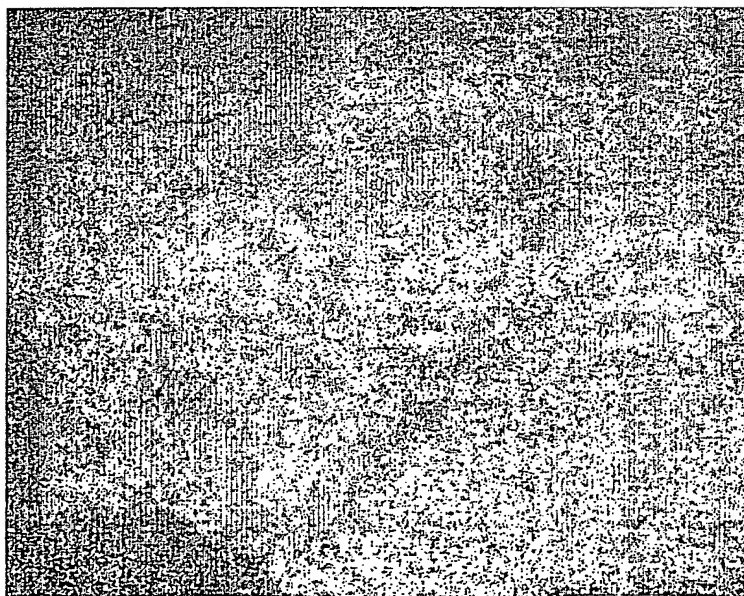


FIGURE 3

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**FIGURE 4**



**FIGURE 5**

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/BR 03/00134-0

## CLASSIFICATION OF SUBJECT MATTER

IPC<sup>7</sup>: A61K 35/78

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC<sup>7</sup>: A61K 35/78

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

WPI, EPODOC, PAJ, MEDLINE

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	JP 2001 122763 A (LION CORP) 8 May 2001 (08.05.01) (abstract). [online] [retrieved on 31.10.2003]. Retrieved from: EPO PAJ Database. <i>abstract.</i>	1-4,6,7,10-13
X	ISOBE T. et al. Antibacterial constituents against Helicobacter pylori of Brazilian medical plant, Pariparoba, Yakugaku zasshi. Journal of the Pharmaceutical Society of Japan, April 2002, Vol. 122, No. 4, pages 291-294, ISSN 0031-6903. Medline-abstract [online] [retrieved on 31 October 2003 (31.10.03)]. Retrieved from: EPOQUE Medline Database, AN NLM11968842. <i>abstract.</i>	4
A	<i>abstract.</i>	1-3,6,7,10-13

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

\* Special categories of cited documents:

„A“ document defining the general state of the art which is not considered to be of particular relevance

„E“ earlier application or patent but published on or after the international filing date

„L“ document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

„O“ document referring to an oral disclosure, use, exhibition or other means

„P“ document published prior to the international filing date but later than the priority date claimed

„T“ later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

„X“ document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

„Y“ document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

„&amp;“ document member of the same patent family

Date of the actual completion of the international search

17 November 2003 (17.11.2003)

Date of mailing of the international search report

3 December 2003 (03.12.2003)

Name and mailing address of the ISA/AT

Austrian Patent Office  
Dresdner Straße 87, A-1200 Vienna

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Authorized officer

WOLF K.

Telephone No. 1/53424/436



## INTERNATIONAL SEARCH REPORT

International application No.

PCT/BR 03/00134-0

## C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	FELZENSZWALB I. et al. Absence of mutagenicity of Potomorphe umbellata and Potomorphe peltata in the Salmonella/mammalian- microsome mutagenicity assay. Brazilian journal of medical and biological research, 1987, Vol. 20, No. 3-4, pages 403-405, ISSN 0100-879X. Medline-abstract [online] [retrieved on 31 October 2003 (31.10.03) ]. Retrieved from: EPOQUE Medline Database, AN NLM3330461. <i>abstract.</i>	4
A	<i>abstract.</i>	1-3,6,7,10-13
A	DE 19933857 A1 (Cognis Deutschland GmbH) 1 February 2001 (01.02.01) --- <i>abstract, claims 1,8.</i>	1-4,6,7,10-13
A	JP 09 208483 A (KAO CORP) 12 August 1997 (12.08.97) (abstract). World Patents Index [online]. London, U.K.: Derwent Publications, Ltd. [retrieved on 31.10.2003]. Retrieved from: Questel/Orbit, Paris, France. DW 9742, Accession No. 97-453955. <i>abstract.</i>	1-4
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# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/BR 03/00134-0

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 11-13  
because they relate to subject matter not required to be searched by this Authority, namely:  
**Remark: Although claims 11-13 concern a method for treatment of the human/animal body by therapy the search has been carried out and based on the alleged effects.**
2. ☐ Claims Nos.:  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

### Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.  
☐ No protest accompanied the payment of additional search fees.

# STATIC VENTING SYSTEM WITH SKYLIGHT

## Information on patent family members

International application No.

PCT/BR 03/00134-0

Patent document cited in search report			Publication date	Patent family member(s)		Publication date
	A				none	
DE	A	19933857	2001-02-01	WO	A 0106996	2001-02-01
JP	A	9208483A			none	
		2				
JP	A	20011227			none	
		63A2				

## PCT REQUEST

Original (for SUBMISSION) - printed on 17.09.2003 03:59:17 PM

VIII-4-1	<p>Declaration: Inventorship (only for the purposes of the designation of the United States of America) Declaration of inventorship (Rules 4.17(iv) and 51 bis.1(a)(iv)) for the purposes of the designation of the United States of America:</p>	<p>I hereby declare that I believe I am the original, first and sole (if only one inventor is listed below) or joint (if more than one inventor is listed below) inventor of the subject matter which is claimed and for which a patent is sought.</p> <p>This declaration is directed to the international application of which it forms a part (if filing declaration with application).</p> <p>I hereby declare that my residence, mailing address, and citizenship are as stated next to my name.</p> <p>I hereby state that I have reviewed and understand the contents of the above-identified international application, including the claims of said application. I have identified in the request of said application, in compliance with PCT Rule 4.10, any claim to foreign priority, and I have identified below, under the heading "Prior Applications," by application number, country or Member of the World Trade Organization, day, month and year of filing, any application for a patent or inventor's certificate filed in a country other than the United States of America, including any PCT international application designating at least one country other than the United States of America, having a filing date before that of the application on which foreign priority is claimed.</p>
VIII-4-1-1	Prior applications:	




PCT/BR 03/00134

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## PCT REQUEST

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PI 0204130-8

		<p>I hereby acknowledge the duty to disclose information that is known by me to be material to patentability as defined by 37 C.F.R. § 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the PCT international filing date of the continuation-in-part application.</p> <p>I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.</p>
VIII-4-1-1-1	Name:	BARROS, Silvia, Berlanga, de, Moraes
VIII-4-1-1-2	Residence: (city and either US State, if applicable, or country)	São Paulo, Brazil
VIII-4-1-1-3	Mailing address:	Rua Caiowáa, 1236/204 São Paulo BR
VIII-4-1-1-4	Citizenship:	
VIII-4-1-1-5	Inventor's Signature: (if not contained in the request, or if declaration is corrected or added under Rule 26ter after the filing of the International application. The signature must be that of the Inventor, not that of the agent)	
VIII-4-1-1-6	Date: (of signature which is not contained in the request, or of the declaration that is corrected or added under Rule 26ter after the filing of the International application)	September 17, 2003

## PCT REQUEST

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VIII-4-1-2-1	Name:	ROPKE, Cristina, Dislich
VIII-4-1-2-2	Residence: (city and either US State, if applicable, or country)	São Paulo, Brazil
VIII-4-1-2-3	Mailing address:	Travessa Francisco Dória de Andrade, 50 São Paulo
VIII-4-1-2-4	Citizenship:	BR
VIII-4-1-2-5	Inventor's Signature: (If not contained in the request, or if declaration is corrected or added under Rule 26ter after the filing of the international application. The signature must be that of the inventor, not that of the agent)	<i>Cristina Dislich Röpke</i>
VIII-4-1-2-6	Date: (of signature which is not contained in the request, or of the declaration that is corrected or added under Rule 26ter after the filing of the international application)	September 17, 2003

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